



King's Research Portal

DOI:

[10.1093/rheumatology/kex315](https://doi.org/10.1093/rheumatology/kex315)

Document Version

Peer reviewed version

[Link to publication record in King's Research Portal](#)

Citation for published version (APA):

Ibrahim, F., Lorente-Cánovas, B., Doré, C. J., Bosworth, A., Ma, M. H., Galloway, J. B., Cope, A. P., Pande, I., Walker, D., & Scott, D. L. (2017). Optimizing treatment with tumour necrosis factor inhibitors in rheumatoid arthritis-a proof of principle and exploratory trial: is dose tapering practical in good responders? *Rheumatology (Oxford, England)*, 56(11), 2004-2014. <https://doi.org/10.1093/rheumatology/kex315>

Citing this paper

Please note that where the full-text provided on King's Research Portal is the Author Accepted Manuscript or Post-Print version this may differ from the final Published version. If citing, it is advised that you check and use the publisher's definitive version for pagination, volume/issue, and date of publication details. And where the final published version is provided on the Research Portal, if citing you are again advised to check the publisher's website for any subsequent corrections.

General rights

Copyright and moral rights for the publications made accessible in the Research Portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognize and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the Research Portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the Research Portal

Take down policy

If you believe that this document breaches copyright please contact librarypure@kcl.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.

Original article

Optimizing treatment with tumour necrosis factor inhibitors in rheumatoid arthritis—a proof of principle and exploratory trial: is dose tapering practical in good responders?

Fowzia Ibrahim¹, Beatriz Lorente-Cánovas¹, Caroline J. Doré², Ailsa Bosworth³, Margaret H. Ma¹, James B. Galloway¹, Andrew P. Cope¹, Ira Pande⁴, David Walker⁵ and David L. Scott¹

Abstract

Objectives. RA patients receiving TNF inhibitors (TNFi) usually maintain their initial doses. The aim of the Optimizing Treatment with Tumour Necrosis Factor Inhibitors in Rheumatoid Arthritis trial was to evaluate whether tapering TNFi doses causes loss of clinical response.

Methods. We enrolled RA patients receiving etanercept or adalimumab and a DMARD with DAS28 under 3.2 for over 3 months. Initially (months 0–6) patients were randomized to control (constant TNFi) or two experimental groups (tapering TNFi by 33 or 66%). Subsequently (months 6–12) control subjects were randomized to taper TNFi by 33 or 66%. Disease flares (DAS28 increasing ≥ 0.6 with at least one additional swollen joint) were the primary outcome.

Results. Two hundred and forty-four patients were screened, 103 randomized and 97 treated. In months 0–6 there were 8/50 (16%) flares in controls, 3/26 (12%) with 33% tapering and 6/21 (29%) with 66% tapering. Multivariate Cox analysis showed time to flare was unchanged with 33% tapering but was reduced with 66% tapering compared with controls (adjusted hazard ratio 2.81, 95% CI: 0.99, 7.94; $P=0.051$). Analysing all tapered patients after controls were re-randomized (months 6–12) showed differences between groups: there were 6/48 (13%) flares with 33% tapering and 14/39 (36%) with 66% tapering. Multivariate Cox analysis showed 66% tapering reduced time to flare (adjusted hazard ratio 3.47, 95% CI: 1.26, 9.58; $P=0.016$).

Conclusion. Tapering TNFi by 33% has no impact on disease flares and appears practical in patients in sustained remission and low disease activity states.

Trial registration: EudraCT, <https://www.clinicaltrialsregister.eu>, 2010-020738-24; ISRCTN registry, <https://www.isrctn.com>, 28955701

Key words: TNF, RA, biologics, tapering, interruption treatment, flare

Rheumatology key messages

- Tapering TNF inhibitors in RA patients by 33% over 6 months did not reduce time to disease flare.
- By 12 months, 45% of RA patients who tapered TNF inhibitor treatment were able to stop it entirely.

¹Academic Department of Rheumatology, Division of Immunology, Infection and Inflammatory Disease, Faculty of Life Sciences and Medicine, King's College London, ²Comprehensive Clinical Trials Unit, University College London, London, ³National Rheumatoid Arthritis Society, Maidenhead, ⁴Rheumatology Department, Nottingham University Hospitals NHS Trust, Nottingham and ⁵Musculoskeletal Services, Freeman Hospital, Newcastle, UK

Submitted 15 November 2016; revised version accepted 12 July 2017

Correspondence to: Fowzia Ibrahim, Academic Department of Rheumatology, Division of Immunology, Infection and Inflammatory Disease, Faculty of Life Sciences and Medicine, King's College London, Weston Education Centre, 10 Cutcombe Road, London SE5 9RJ, UK.
E-mail: fowzia.ibrahim@kcl.ac.uk

Introduction

Trials and observational studies underpinning the regulatory approval of TNF inhibitors (TNFi) for treating RA patients focus on initial efficacy and long-term safety [1–3]. What to do after achieving disease control is another question. Currently doses of TNFi effective in inducing responses in active RA are continued to maintain control, despite limited evidence they are needed. Maintaining disease control maintained with lower doses of TNFi should increase their cost-effectiveness.

The Optimizing Treatment with TNFi in RA (OPTTIRA) trial was designed when there was limited information about tapering TNF inhibitor in RA patients with good treatment responses. Subsequently several observational studies and trials evaluated tapering. Systematic reviews of reports before 2015 [4–7] found evidence supporting TNF tapering. The Cochrane systematic review, which focused exclusively on trials [7], found tapering gave similar outcomes to continuing standard doses of adalimumab and etanercept; complete discontinuation increased flares. Another seven trials [8–14] were published after these systematic reviews were completed. One trial [12] concluded disease activity guided adalimumab or etanercept dose reductions were non-inferior to usual care for major flares; dose reduction or stopping was possible in two-thirds of patients. Overall the evidence suggests tapering does not substantially increase flares; stopping completely may do so. Recent observational studies support dose reduction [15–17]. Reviews by Schett *et al.* [18] and Edwards *et al.* [19] highlighted the importance of TNF inhibitor tapering and minimizing other long-term DMARDs. However, there remain uncertainties about which patients should have their TNFi tapered and whether all tapering regimens are similar.

OPTTIRA is a randomized trial evaluating two tapering regimens in RA. It recruited patients showing EULAR good responses to TNFi [20] and compared tapering with continuing standard doses. Tapering regimens reduced doses to one-third and two-thirds initial response induction doses of adalimumab or etanercept over 6 months. OPTTIRA also examined subsequently stopping TNFi completely. It used time to flare to assess the effects of TNF inhibitor tapering. Flares occurred when the DAS28 joints was over 3.2 and increased by 0.6 or more.

Methods

Design

OPTTIRA was an open label, 6-month multicentre proof of principle trial with a subsequent 6-month exploratory phase for patients who completed the initial trial. OPTTIRA enrolled RA patients achieving good responses [20] with low disease activity or remission taking standard TNFi doses and receiving one or more DMARDs.

Participants

Patients receiving TNFi had met existing English criteria from the National Institute for Health and Clinical

Excellence for these agents. The criteria have changed with time; they included failing to respond to MTX and another DMARD [21]. These criteria meant all patients had established RA. Patients had also achieved sustained good responses with DAS28 scores of ≤ 3.2 without increases of > 0.6 during the previous 3 months.

Interventions

Patients were taking etanercept or adalimumab; their existing TNFi were the trial investigational medicinal products.

The proof of principle trial (months 0–6) compared three groups: experimental group 1: TNF inhibitor tapered by 33% initial dose; experimental group 2: TNF inhibitor tapered by 66% initial dose; control group: continued standard doses. Supplementary Tables S1 and S2, available at *Rheumatology* Online, show reducing etanercept and adalimumab dosing schedules.

In the exploratory phase (months 7–12) patients in experimental groups increased times between injections until they stopped. Patients in the control group were further randomized into two groups: control group A had TNFi tapered by 33% initial dose; control group B had TNFi tapered by 66% initial dose. Supplementary Tables S3 and S4, available at *Rheumatology* Online, show the dosing schedules. The tapering schedules reflected standard dosing regimens related to the half-lives of the drugs.

Primary outcome

The primary outcome was time to first flare, defined as an increase in DAS28 scores ≥ 0.6 resulting in a DAS28 > 3.2 together with an increase in the swollen joint count; both had to be present on two occasions at least 1 week apart. An increase in DAS28 score ≥ 1.2 resulting in DAS28 > 3.2 was defined as flare irrespective of changes in swollen joints. These criteria reflect the subsequently developed DAS28 flare definitions proposed by OMERACT [22] and supported by the Cochrane group [7]. Patients were assessed 3 monthly and telephoned by their Research Nurse monthly to check their disease control. Patients who considered they were experiencing a flare were seen urgently (within 2 weeks).

Secondary outcomes

HAQ, EuroQol 5-dimension scale (EQ5D-3L), Medical Outcomes Study 36-Item Short Form Health Survey (SF-36) and Functional Assessment of Chronic Illness Therapy were assessed at 0, 3, 6 and 12 months. X-rays of the hands (including wrists) and feet were taken at 0, 6 and 12 months with digitized X-rays read by an experienced observer (D.L.S.) blinded to treatment using modified Larsen scores. Every 3 months disease activity assessments recorded tender and swollen joint counts (28 joints), ESR, patients' global assessments of disease activity (100 mm visual analogue scale) and DAS for 28 joints together with details of medication and adverse effects. An anonymized electronic data capture system

collected clinical data except X-rays (<http://www.medsci-net.net>).

Sample size

The TEMPO trial showed 15% of patients on etanercept with MTX withdrew annually (flares and other problems) [23, 24]. The BSR Biologics Register [25] showed 12% of patients withdraw annually from TNFi. These results suggested 12–15% of patients taking TNFi flared annually; during the 6 months proof of principle trial the likelihood of flare was 7.5%. Studying 30 patients in one tapering group would show a significant difference at the 5% level with 80% power if over 42% of patients flared; such a difference would mean tapering was not clinically useful. Therefore a proof of principle study of 30 patients in each group should provide sufficient information to reject the concept if it is clearly ineffective, as well as giving enough data to design larger trials if needed. We allowed for 10% of patients not continuing in the trial and therefore aimed to recruit 33 patients in each group with a total study size of 99 patients.

The randomization algorithm considered patients in both proof of principle trial and the exploratory study taking different biologics. Consequently the randomization ratio generated was 1:1:2; we recruited twice as many to the control group than the initial tapering groups in the proof of principle trial.

Randomization

Potentially eligible patients were screened and reasons for non-entry recorded. The electronic data capture system (MedSciNet) randomized patients using minimization into experimental and control groups, stratified by TNF inhibitor (etanercept or adalimumab) using randomly permuted blocks. Recruiting staff were blinded to allocation sequences.

Blinding

OPTTIRA was unblinded for assessors because it used treatment-specific algorithms with the adjustment of multiple dosing intervals of the drugs. Disability and quality of life were assessed by patients and X-ray reading was blinded.

Statistical methods

Randomized patients accepting their allocated treatment were analysed on an intention-to-treat basis. Data management and analyses used Stata (version 14.0, StataCorp, College Station, TX, USA). Baseline characteristics were summarized by randomization group as means and standard deviations (normally distributed variables), medians and interquartile ranges (non-normally distributed variables) and frequencies and percentages (categorical variables). Serious adverse event rates in each arm were compared with controls.

Separate analysis was performed for the proof of principle phase (0–6 months) and the exploratory phase (6–12 months). The primary outcome was time to flare (months) for patients randomized to control, 33% taper or 66% taper, defined as the time from study entry to

the first flare. Patients without flares, who withdrew or were lost to follow-up, were censored at the time of their last visit. Eight out of 97 patients were lost to follow-up (supplementary Fig. S1, available at *Rheumatology* Online). Kaplan–Meier curves and the log-rank test compared randomized groups. Survival analysis analysed time to flare. The validity of the assumption of proportionality required for Cox regression was investigated graphically (using Nelson–Aalen plots) as well as using Schoenfeld residuals in the final Cox models.

Secondary outcomes were analysed using mixed models to estimate treatment effects including baseline values as covariates. Working correlation matrices were unstructured, which is not unduly restrictive given that measurements are only taken at three time points. The sandwich estimator of covariance matrix was used in order to obtain appropriate (consistent) estimates of precision. All P-values were two sided.

Ethical review

The North West London Research Ethics Committee approved OPTTIRA (REC Ref: 10/H0720/69). All enrolled patients gave written informed consent. The trial was registered with the UK Clinical Research Network and other relevant organizations (EudraCT number: 2010-020738-24; ISRCTN: 28955701).

Results

Patients and treatments

Recruitment

Between April 2011 and June 2013, 244 patients were screened, 103 randomized and 97 accepted their allocated treatment (supplementary Fig. S1, available at *Rheumatology* Online).

Treatment at baseline

All patients received disease modifying drugs: 82 had DMARD monotherapy; 15 received combination DMARDs. The DMARDs comprised MTX (81 patients), HCQ (18), SSZ (11) and LEF (4). All patients received etanercept (43 patients) or adalimumab (54). Two patients were taking prednisolone. The mean baseline treatment duration was 5.8 years (range 0.5–15.1 years).

Proof of principle phase

Of the patients, 50/97 were randomized to continue biologic at unchanged dosages, 26/97 to taper biologics by 33% and 21/97 to taper by 66%; 74/97 patients completed 6 months' treatment, 8/97 were lost to follow-up and 15/97 discontinued tapering but were followed up; 13/97 patients stopped treatment for flares, 1/97 stopped for drug toxicity and 9/97 stopped for other reasons including 1/97 for disease progression and 5/97 at the patients' own request.

Exploratory phase

Of the patients randomized to continue biologics at unchanged doses who completed the 6-month proof of

principle trial, 40/50 were re-randomized in the exploratory phase: 22/40 tapered by 33% and 18/40 tapered by 66%; 32/40 patients completed 6 months' treatment and 8/40 patients discontinued tapering but were followed up; 5/40 stopped treatment for flares, 2/40 had disease progression and 1/40 stopped at the patients' own request.

The 34 patients who had tapered for 6 months and continued on the treatment schedule in the proof of principle trial tapered further until their biologics were completely stopped by 6 months; 21/34 completed 6 months' treatment, 2/34 were lost to follow-up and 11/34 discontinued tapering but were followed up. All 13/34 patients stopping treatment did so because of flares.

Baseline data and numbers analysed

Demographic and disease assessments were similar in all patient groups (Table 1). The patients had established RA (median disease duration 11.3 years) with low DAS28-ESR (median 2.0) and low HAQ scores (median 0.50). All treated patients were analysed in the proof of principle trial.

Primary outcome

Proof of principle phase

Seventeen patients flared comprising 8/50 (16%) control patients, 3/26 (12%) patients tapering by 33% and 6/21 (29%) patients tapering by 66% (Table 2). Univariate and multivariate Cox analysis of the intention to treat population (Table 2) showed no evidence 33% tapering reduced time to flare (adjusted hazard ratio 0.87, 95% CI: 0.22, 3.88; $P=0.835$), but 66% tapering significantly reduced time to flare compared with controls (adjusted hazard ratio 2.81, 95% CI: 0.99, 7.94; $P=0.051$). Figure 1 shows Kaplan-Meier survival curves comparing groups.

Exploratory phase—re-randomized patients

Eleven patients flared: 3/22 (14%) patients tapering by 33% and 8/18 (44%) tapering by 66% (Table 2). Univariate and multivariate Cox analysis showed 66% tapering significantly increased the risk of flare compared with 33% tapering (adjusted hazard ratio 5.10, 95% CI: 1.81, 21.95; $P=0.029$).

Combined tapering groups. Results from re-randomized initial controls were combined with the tapering groups in the proof of principle trial to provide a detailed comparison of time to flare in the two tapering regimens over 6 months. Six of forty-eight (13%) patients tapering by 33% flared compared with 14/39 (36%) patients tapering by 66% (Table 2). Multivariate Cox analysis showed 66% tapering significantly increased the risk of flare compared with 33% tapering (adjusted hazard ratio 3.47, 95% CI: 1.26, 9.58; $P=0.016$).

Secondary outcomes in proof of principle trial

Longitudinal analysis of changes within the first 6 months in all treated patients (Table 3) showed little evidence that 33% tapering had any impact on clinical or functional assessments. However, 66% tapering led to significant worsening in tender and swollen joint counts, C-reactive protein levels and ESR and DAS28 ESR but not DAS28

CRP scores. An interesting finding is that patient related outcome measures including patient global score, HAQ, EQ5D-3L, visual analogue scale (VAS) pain and functional assessment of chronic illness therapy fatigue scores were unaffected by tapering, including tapering by 66%. Assessor global scores and Larsen scores also did not differ between groups. Details of secondary outcomes are shown in supplementary Table S5, available at *Rheumatology Online*.

Impact of flares in proof of principle phase

The effects of flares, irrespective of treatment group, on 6-month outcomes were assessed in linear regression models adjusted for age, gender and disease duration. DAS28 scores were higher in patients who flared (0.85, 95% CI: 0.44, 1.26; $P<0.001$) and EQ5D scores were lower (-0.12 , 95% CI: -0.23 , -0.02 ; $P=0.022$), but HAQ scores were unaffected (0.92, 95% CI: -0.19 , 0.22; $P=0.87$).

Discontinuing TNF inhibitor

Thirty-four patients who tapered biologics in the proof of principle trial (21 with 33% tapering; 13 with 66% tapering) further tapered treatment until they stopped TNF inhibitor over 6 months in the exploratory study. Thirteen of thirty-four (38%) flared and 21/34 (62%) did not flare. Consequently 21/47 (45%) patients who started tapering in the proof of principle trial stopped TNF inhibitor by 12 months; 21/47 (45%) had flared and 5/47 (11%) had stopped tapering for other reasons. The 21 patients who stopped TNF inhibitor without flaring had initial DAS28 score of 1.51 (95% CI: 1.14, 2.65) and 12 month DAS28 scores of 2.27 (95% CI: 1.71, 3.98). Their initial HAQ was 0.65 (95% CI: 0.33, 0.99) and the 12 month HAQ 0.75 (95% CI: 0.42, 1.17).

Flares in patient subgroups

The flare rate for patients receiving adalimumab, 18/54 (33%), was significantly lower ($\chi^2=3.99$, degrees of freedom (DF)=1, $P=0.050$) than for patients receiving etanercept, 23/43 (53%).

At baseline 74 patients were in remission and had DAS28 scores under 2.6; 27/74 (36%) of these patients flared. There were 23 patients with low disease activity (DAS28 score 2.6–3.2) and 14/23 (61%) of these flared. The risk of flare was therefore significantly higher for patients with low disease activity at baseline ($\chi^2=4.3$, DF=1, $P=0.039$) than for those in remission.

Adverse events

Overall there were 443 adverse events; 76/97 (78%) patients had one or more events; 47/97 (48%) patients had an adverse event in the first 6 months and 29/89 (33%) in the second 6 months (Table 4); 4/97 (4%) of patients had a serious adverse event, 3 in the first 6 months and 1 in the second 6 months; 220/443 (50%) of adverse events involved the musculoskeletal system and only 223/443 (50%) involved other systems. No relationships were identified between treatment tapering and adverse events.

TABLE 1 Baseline demographic and clinical characteristics

	33% tapering n = 26	66% tapering n = 21	Controls n = 50	Total n = 97
Demographic variables				
Age, mean (s.d.), years	59 (11)	58 (9)	56 (12)	57 (11)
Height, mean (s.d.), m	1.66 (0.08)	1.67 (0.08)	1.66 (0.09)	1.66 (0.08)
Weight, mean (s.d.), kg	74.8 (15.6)	70.1 (14.4)	74.3 (16.1)	73.5 (15.5)
BMI, kg/m ²	26.9 (21.9–31.6)	24.5 (22.6–27.8)	25.3 (23.1–29.7)	25.4 (22.6–29.4)
Disease duration, years	11.2 (6.2–19.0)	10.6 (7.3–15.9)	11.9 (7.3–16.7)	11.3 (7.3–16.7)
Female gender, n (%)	19 (73)	15 (71)	38 (76)	72 (74)
Smoking status, n (%)				
Ex	9 (43)	7 (41)	16 (35)	32 (38)
Current	2 (9)	3 (18)	7 (15)	12 (14)
Clinical variables				
Tender joint counts, 28 joints	0 (0–0)	0 (0–0)	0 (0–1.00)	0 (0–1.00)
Swollen joint counts, 28 joints	0 (0–0)	0 (0–0)	0 (0–0)	0 (0–0)
Tender joint counts, 68 joints	0 (0–1.00)	0 (0–2.00)	0 (0–2.00)	0 (0–2.00)
Swollen joint counts, 66 joints	0 (0–0)	0 (0–0)	0 (0–0)	0 (0–0)
ESR, mm/h	6 (4–24)	8 (4–19)	9 (5–20)	8 (5–19)
CRP, mg/l	5 (2–6)	4 (2–5)	5 (2–7)	5 (2–6)
Assessor global rating, mm	3 (0–10)	4 (1–10)	3 (1–9)	3 (1–10)
Patient global assessment, mm	9 (1–15)	4 (1–16)	5 (2–16)	5 (1–16)
DAS28 ESR	1.7 (1.1–2.6)	1.9 (1.3–2.6)	2.1 (1.4–2.6)	2.0 (1.3–2.6)
DAS28 CRP	2.3 (2.0–2.5)	2.2 (1.7–2.5)	2.1 (1.9–2.5)	2.2 (1.9–2.5)
HAQ	0.75 (0.13–1.38)	0.38 (0.0–0.88)	0.50 (0.13–1.50)	0.50 (0.13–1.38)
EQ5D-3L	0.79 (0.66–1.00)	0.80 (0.69–1.00)	0.74 (0.59–1.00)	0.76 (0.66–1.00)
Pain visual analogue score, mm	6 (1–10)	8 (1–19)	5 (0–25)	5 (1–19)
FACIT Fatigue Scale	40 (31–45)	41 (35–46)	42 (36–46)	41 (35–46)
Larsen score	33 (12–76)	34 (17–63)	66 (29–89)	51 (16–82)
SF-36				
PCS	43 (38–51)	46 (37–51)	44 (32–52)	45 (34–52)
MCS	51 (43–58)	57 (48–60)	58 (52–61)	57 (49–60)

Values shown as median (interquartile range) unless otherwise stated. EQ5D-3L: EuroQol 5-dimension scale; FACIT: Functional Assessment of Chronic Illness Therapy; MCS: mental health summary score; PCS: physical health summary score; SF-36: 36-Item Short Form Health Survey.

TABLE 2 Flare rates and univariate and multivariate cox analyses in intention to treat analysis and exploratory studies

			Unadjusted		Adjusted ^a	
	Flares, n (%)		HR (95% CI)	P-value	HR (95% CI)	P-value
Intention to treat analysis (n = 97)	Control	8/50 (16)	Reference		Reference	
	33% tapering	3/26 (12)	0.90 (0.23, 3.48)	0.873	0.87 (0.22, 3.88)	0.835
	66% tapering	6/21 (29)	2.52 (0.85, 7.48)	0.097	2.81 (0.99, 7.94)	0.051
Exploratory study (n = 40)	33% tapering	3/22 (14)	Reference		Reference	
	66% tapering	8/18 (44)	4.16 (1.08, 15.99)	0.038	5.10 (1.18, 21.95)	0.029
Combined tapering (n = 87)	33% tapering	6/48 (13)	Reference		Reference	
	66% tapering	14/39 (36)	3.29 (1.26, 8.63)	0.015	3.47 (1.26, 9.58)	0.016

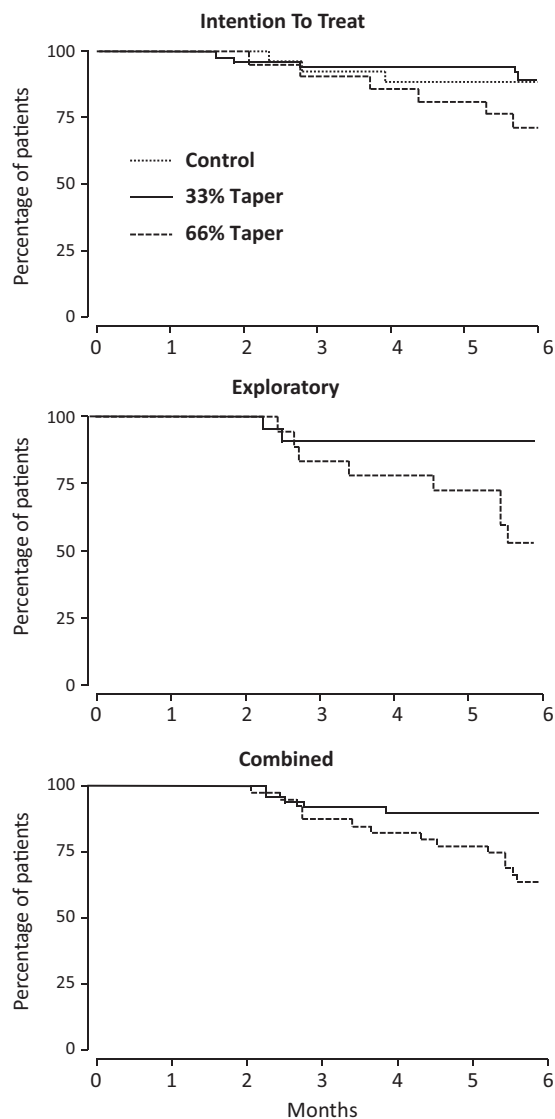
^aThe multivariate model was adjusted for age at registration, gender and disease duration.

Discussion

OPTTIRA shows reducing TNF inhibitor doses by one-third in RA patients with stable low disease activity or remission also taking DMARDs has no impact on disease

activity or the frequency of flares. However, the limited number of patients included in each arm restricts the importance of its finding for clinical practice. Most patients were in remission; there was some evidence patients with low disease activity scores had more flares. Greater

Fig. 1 Kaplan–Meier curves for intention to treat analysis, exploratory and combined tapering groups



reductions in TNF inhibitor doses led to more flares and higher disease activity levels but had no impact on disability. TNFi could be stopped in many patients without major negative impacts upon their disease.

The balance of evidence from OPTTIRA and previous trials and observational studies of tapering and stopping TNFi is that modest dose reduction is possible in RA patients with good responses to TNFi who remain on conventional DMARDs. Indeed there is little evidence favouring maintaining standard doses in such patients. OPTTIRA showed that reducing biologic doses by one-third had no impact on clinical, functional or health status assessments and did not result in more flares. TNFi are expensive with uncertainties about their overall cost-effectiveness [26, 27] and in patients with good

responses tapering appears less expensive than maintaining full doses without major negative impacts.

The evidence favouring major reductions in TNF inhibitor dose, such as reducing by two-thirds, is less clear-cut. OPTTIRA, other trials and observational studies all found this gives more flares and worse clinical and health status. However, the negative impacts are modest and in some patients treatment can be stopped without obvious disadvantages. Considering stopping treatment may be reasonable in selected cases.

Reductions in TNF inhibitor doses in stable RA have not been approved by regulatory agencies and therefore organizations like the National Institute for Health and Clinical Excellence cannot recommend such approaches [28]. Nevertheless, individual clinicians may decide, based on overall assessments of risks and benefits, to offer dose reductions to some patients. The impact of tapering TNFi is broadly similar to reducing the doses of conventional DMARDs; reducing MTX doses can be achieved in some patients but stopping treatment increases flares. Defining patients likely to flare is an important future research goal.

Adverse events were common in OPTTIRA and often involved the musculoskeletal system. Only four were serious events and these were unrelated to tapering or stopping treatment. The serious adverse event rate was similar to that in the systematic review of tapering by van Herwaarden *et al.* [7] who reported 5% of patients had serious adverse events. Such adverse event rates are likely to be commonplace in treated RA patients.

TNFi are expensive, and while using effective biosimilars [29] will reduce costs, such biologic treatments will never become inexpensive. Consequently the clinical and the economic benefits of good responders remaining on standard doses of TNFi will diverge. As the disadvantages of tapering and trying to stop TNFi are relatively small, clinicians and healthcare funders may consider it preferable to explore reducing and stopping treatment when patients achieve good disease control. In early RA, using biologics initially may result in sustained biologic-free and drug-free remissions [30, 31]; OPTTIRA and similar trials have not evaluated this possibility. Nevertheless it may be important because achieving sustained deep remission may be of critical consequence in early disease.

Our trial had several strengths. It involved a range of RA patients across multiple English centres, making its findings likely to be generalizable. It evaluated two tapering regimens and stopping treatment, ensuring several strategies for reducing TNF inhibitor doses. It had several weaknesses. First, it was relatively small and enrolled only 97 patients. Its lack of power precludes robust conclusions in secondary analyses evaluating the impact of tapering on HAQ within the 66% tapering group. Secondly, it was relatively short and assessors were not blinded, in keeping with other pragmatic trials in the area. Thirdly, it did not look at all responders to TNFi, only those who had sustained low disease activity or remission; patients who almost achieved low disease activity may respond similarly to those enrolled in the trial. Fourthly, it only studied two TNFi, and may not apply to other TNFi.

TABLE 3 Secondary outcomes for the effect of randomized treatment arm in longitudinal analysis for first 6 months

Outcome	33% tapering			66% tapering		
	Unadjusted coefficient (95% CI)	P-value	Adjusted coefficient ^a (95% CI)	Unadjusted coefficient (95% CI)	P-value	Adjusted coefficient ^a (95% CI)
DAS28-ESR	0.13 (−0.20, 0.47)	0.438	0.06 (−0.25, 0.38)	0.44 (0.07, 0.80)	<0.02	0.36 (0.02, 0.70)
DAS28-CRP	−0.05 (−0.27, 0.18)	0.671	−0.05 (−0.27, 0.18)	0.17 (−0.08, 0.41)	0.188	0.18 (−0.07, 0.42)
Tender joint counts	0.07 (−0.76, 0.91)	0.864	0.05 (−0.78, 0.88)	1.34 (0.42, 2.25)	<0.01	1.32 (0.42, 2.22)
Swollen joint counts	−0.17 (−0.71, 0.37)	0.530	−0.21 (−0.74, 0.32)	0.97 (0.39, 1.56)	0.001	0.88 (0.30, 1.46)
ESR	−0.97 (−5.06, 3.11)	0.641	−1.07 (−4.15, 2.00)	−3.47 (−7.89, 0.96)	0.124	−3.87 (−7.21, −0.53)
CRP	−2.60 (−5.50, 0.30)	0.079	−2.86 (−5.67, −0.05)	−4.34 (−7.49, −1.18)	0.01	−4.28 (−7.34, −1.21)
Assessor global	0.86 (−5.15, 6.86)	0.779	0.72 (−5.30, 6.73)	5.23 (−1.27, 11.73)	0.115	5.11 (−1.40, 11.63)
Patient global	2.64 (−3.32, 8.59)	0.385	3.00 (−2.73, 8.74)	4.82 (−1.62, 11.27)	0.142	5.34 (−0.87, 11.55)
HAQ	0.04 (−0.10, 0.17)	0.591	0.03 (−0.10, 0.15)	0.15 (0.01, 0.30)	<0.05	0.13 (−0.01, 0.27)
EQ5D-3L	−0.04 (−0.12, 0.03)	0.249	−0.03 (−0.10, 0.04)	−0.05 (−0.13, 0.03)	0.248	−0.04 (−0.11, 0.04)
VAS pain	6.62 (−1.83, 15.06)	0.125	5.10 (−2.49, 12.69)	3.85 (−5.30, 12.99)	0.410	0.79 (−7.55, 9.13)
FACIT fatigue	−0.35 (−2.93, 2.24)	0.793	−0.58 (−3.01, 1.85)	1.41 (−1.38, 4.20)	0.321	1.03 (−1.60, 3.65)
Larsen scores	−0.21 (−1.69, 1.27)	0.780	0.13 (−1.39, 1.64)	0.35 (−1.25, 1.96)	0.867	0.63 (−0.98, 2.25)

^aThe model was adjusted for age at registration, gender and disease duration. EQ5D-3L: EuroQol 5-dimension scale; FACIT: Functional Assessment of Chronic Illness Therapy; VAS: visual analogue scale.

TABLE 4 Serious and adverse events by body system and intervention group

Body system	Proof of principle phase (0-6 months)						Exploratory phase (6-12 months)						Re-randomized control group to								
	Tapered 33%, n = 15 (96 episodes)			Tapered 66%, n = 13 (71 episodes)			Controls, n = 19 (120 episodes)			Tapering 33% until stop, n = 9 (54 episodes)			Tapering 66% until stop, n = 4 (28 episodes)			Tapered 33%, n = 10 (43 episodes)			Tapered 66%, n = 6 (35 episodes)		
	Serious	All		Serious	All		Serious	All		Serious	All		Serious	All		Serious	All		Serious	All	
Cardiovascular		3			1			3		0			0			2					0
Digestive		13			3			12		2			0			6					7
ENT		5			1			5		4			2			5					7
Endocrine/metabolic		0			0			0		0			0			0					0
Genitourinary	1	1			2			3		0			0			2					0
Haematological		0		1	1			6		1			0			0					0
Mental		0			0			1		0			2			0					0
Musculoskeletal		51			37			36		45			18			19					5
Nervous system		4			1			4		0			0			0					14
Ophthalmological		0			0			4		0			0			0					0
Respiratory		8			8			33		2			1			8					7
Skin	1	11			0			13		0			5			0			1		0

Seventy-six patients had an adverse event; of those, 47 patients had an AE in the first 6 months and 29 in the next 6 months.

Fifthly, it did not consider sustained flares though there is some evidence [32] these are more important than transient flares, nor did it assess flares from patients' perspectives, which recent research has highlighted as being relevant [33–35]. Sixthly, many patients did not wish to participate in OPTTIRA; our non-participation rate was substantially higher than some trials like that reported by Moghadam *et al.* [11] but lower than others [8]. Similar high non-participation rates have occurred in other RA trials from our unit [36, 37]. This difference may represent different ways of collecting data about screening or national differences in patients' views on trials. However, caution is needed when considering our results' generalizability. Finally, only limited information was available on radiological outcomes.

An issue in RA treatment, particularly relevant tapering biologics, is identifying patient sub-groups likely to respond well to the treatment strategy. Analysis of predictors in the BeSt trial [38] showed Anti-citrullinated protein antibody (ACPA) was an important indicator of flares after stopping infliximab. The impact of ACPA status on response to tapering was also reported by Haschka *et al.* [10]. More recently Rech *et al.* [39] showed combining ACPA status with multibio-marker disease activity assessment predicted relapses in over 80% of patients. An alternative is using US assessments to predict response to tapering [40, 41]. There is evidence in our trial that tapering may be best in patients achieving deep remissions; further work is needed to clarify this possibility.

The most effective and cost-effective strategy for using biologics in RA patients remains uncertain. As current biologics are not curative in established RA patients, the rationale for their indefinite use in treatment responders is debatable. OPTTIRA shows modest tapering has no negative clinical impacts. We consider there is sufficient evidence for clinicians to reduce TNF inhibitor doses in some treatment responders. This approach may be more cost-effective than their continued use though it risks short-term disadvantages as flares are a burden to patients. Its impacts on disease status and healthcare costs need further evaluation, particularly as we found no evidence that tapering worsened patient-assessed outcomes.

Acknowledgements

We are grateful for the many colleagues and collaborators who contributed to the OPTTIRA trial. *Trial Office Staff:* Nestor Salazar, Amaka Ejibe, Rebecca Brendell, Dr Lisa Ann Clutterbuck, Dr Anna Kowalczyk, Beverley White-Alao, Janice Jimenez. *Trial Steering Committee:* Professor David G. I. Scott, Caroline J. Doré, Dr Andrew Keat, Dr Brian Tom, Dr David Collins, Professor Simon Bowman, Ms Jacqui Deakin. *Data Monitoring and Ethics Committee:* Dr Chris Deighton (Chair), Dr Richard Watts, Professor Alex Macgregor. *Principal Investigators:* Dr Anurag Bharadwaj (Basildon University Hospital), Professor Andrew Cope (Guy's Hospital), Dr Ira Pande (Queens Medical Centre), Dr Sarah Levy (Croydon

University Hospital), Dr David Walker (Freeman Hospital), Dr Michael Plant (James Cook University Hospital), Dr Stuart Kyle (North Devon Hospital), Dr Sarah Medley (Orpington Hospital), Dr Selwyn Richards (Poole Hospital), Dr Clive Kelly (Queen Elizabeth Gateshead Hospital), Dr Gerald Coakley (Queen Elizabeth Hospital, Woolwich) Dr Nap Cheung (Queen Mary's Hospital), Dr James Maxwell (Rotherham General Hospital), Professor Anthony Woolf (Royal Cornwall Hospital), Dr Richard Haigh (Royal Devon and Exeter Hospital), Dr Karen Walker-Bone (Royal Sussex County Hospital), Dr David Collins (The Great Western Hospital), Dr Kirsten Mackay (Torbay Hospital), Professor Gabrielle Kingsley (University Hospital Lewisham). *Patient Involvement:* Patients were involved in the design of the trial and a co-applicant on the grant. One this patient was on the trial steering committee throughout the trial.

Funding: The trial was funded by Arthritis Research UK (grant reference number 18813).

Disclosure statement: J.B.G. has received honoraria for speaking from UCB, Pfizer, Celgene and Bristol-Myers Squibb. All other authors have declared no conflicts of interest.

Supplementary data

Supplementary data are available at *Rheumatology Online*.

References

- de La Forest Divonne M, Gottenberg JE, Salliot C. Safety of biologic DMARDs in RA patients in real life: A systematic literature review and meta-analyses of biologic registers. *Joint Bone Spine* 2017;84:133–40.
- Hazlewood GS, Barnabe C, Tomlinson G *et al.* Methotrexate monotherapy and methotrexate combination therapy with traditional and biologic disease modifying antirheumatic drugs for rheumatoid arthritis: abridged Cochrane systematic review and network meta-analysis. *BMJ* 2016;353:i1777.
- Orme ME, Macgilchrist KS, Mitchell S, Spurdin D, Bird A. Systematic review and network meta-analysis of combination and monotherapy treatments in disease-modifying antirheumatic drug-experienced patients with rheumatoid arthritis: analysis of American College of Rheumatology criteria scores 20, 50, and 70. *Biologics* 2012;6:429–64.
- Galvao TF, Zimmermann IR, da Mota LM, Silva MT, Pereira MG. Withdrawal of biologic agents in rheumatoid arthritis: a systematic review and meta-analysis. *Clin Rheumatol* 2016;35:1659–68.
- Kuijper TM, Lamers-Karnebeek FBG, Jacobs JWG, Hazes JM, Luime JJ. Flare rate in patients with rheumatoid arthritis in low disease activity or remission when tapering or stopping synthetic or biologic DMARD: a systematic review. *J Rheumatol* 2015;42:2012–22.

- 6 Navarro-Millan I, Sattui SE, Curtis JR. Systematic review of tumor necrosis factor inhibitor discontinuation studies in rheumatoid arthritis. *Clin Ther* 2013;35:1850–61.e1.
- 7 van Herwaarden N, den Broeder AA, Jacobs W *et al*. Down-titration and discontinuation strategies of tumor necrosis factor-blocking agents for rheumatoid arthritis in patients with low disease activity. *Cochrane Database Syst Rev* 2014;(9):CD010455.
- 8 Chatzidionysiou K, Turesson C, Telesman A *et al*. A multicentre, randomised, controlled, open-label pilot study on the feasibility of discontinuation of adalimumab in established patients with rheumatoid arthritis in stable clinical remission. *RMD Open* 2016;2:e000133.
- 9 Fautrel B, Pham T, Alfaiaite T *et al*. Step-down strategy of spacing TNF-blocker injections for established rheumatoid arthritis in remission: results of the multicentre non-inferiority randomised open-label controlled trial (STRASS: Spacing of TNF-blocker injections in Rheumatoid Arthritis Study). *Ann Rheum Dis* 2016;75:59–67.
- 10 Haschka J, Englbrecht M, Hueber AJ *et al*. Relapse rates in patients with rheumatoid arthritis in stable remission tapering or stopping antirheumatic therapy: interim results from the prospective randomised controlled RETRO study. *Ann Rheum Dis* 2016;75:45–51.
- 11 Moghadam MG, Vonkeman HE, ten Klooster PM *et al*. Stopping tumor necrosis factor inhibitor treatment in patients with established rheumatoid arthritis in remission or with stable low disease activity. *Arthritis Rheumatol* 2016;68:1810–7.
- 12 van Herwaarden N, van der Maas A, Minten MJ *et al*. Disease activity guided dose reduction and withdrawal of adalimumab or etanercept compared with usual care in rheumatoid arthritis: open label, randomised controlled, non-inferiority trial. *BMJ* 2015;350:h1389.
- 13 van Vollenhoven RF, Ostergaard M, Leirisalo-Repo M *et al*. Full dose, reduced dose or discontinuation of etanercept in rheumatoid arthritis. *Ann Rheum Dis* 2016;75:52–8.
- 14 Yamanaka H, Nagaoka S, Lee SK *et al*. Discontinuation of etanercept after achievement of sustained remission in patients with rheumatoid arthritis who initially had moderate disease activity—results from the ENCOURAGE study, a prospective, international, multicenter randomized study. *Mod Rheumatol* 2016;26:651–61.
- 15 Tanaka Y, Hirata S, Kubo S *et al*. Discontinuation of adalimumab after achieving remission in patients with established rheumatoid arthritis: 1-year outcome of the HONOR study. *Ann Rheum Dis* 2015;74:389–95.
- 16 Yoshida K, Kishimoto M, Radner H *et al*. Low rates of biologic-free clinical disease activity index remission maintenance after biologic disease-modifying antirheumatic drug discontinuation while in remission in a Japanese multicentre rheumatoid arthritis registry. *Rheumatology* 2016;55:286–90.
- 17 Yoshida K, Radner H, Mjaavatten MD *et al*. Incidence and predictors of biological antirheumatic drug discontinuation attempts among patients with rheumatoid arthritis in remission: A CORRONA and NinJa Collaborative Cohort Study. *J Rheumatol* 2015;42:2238–46.
- 18 Schett G, Emery P, Tanaka Y *et al*. Tapering biologic and conventional DMARD therapy in rheumatoid arthritis: current evidence and future directions. *Ann Rheum Dis* 2016;75:1428–37.
- 19 Edwards CJ, Fautrel B, Schulze-Koops H, Huizinga TW, Kruger K. Dosing down with biologic therapies: a systematic review and clinicians' perspective. *Rheumatology* 2017;56:1847–56.
- 20 van Gestel AM, Haagsma CJ, Riel PLCM. Validation of rheumatoid arthritis improvement criteria that include simplified joint counts. *Arthritis Rheum* 1998;41:1845–50.
- 21 NICE. Adalimumab, etanercept, infliximab, certolizumab pegol, golimumab, tocilizumab and abatacept for rheumatoid arthritis not previously treated with DMARDs or after conventional DMARDs only have failed. 2016. <https://www.nice.org.uk/guidance/ta375>.
- 22 van der Maas A, Lie E, Christensen R *et al*. Construct and criterion validity of several proposed DAS28-based rheumatoid arthritis flare criteria: an OMERACT cohort validation study. *Ann Rheum Dis* 2013;72:1800–5.
- 23 van der Heijde D, Klareskog L, Landewe R *et al*. Disease remission and sustained halting of radiographic progression with combination etanercept and methotrexate in patients with rheumatoid arthritis. *Arthritis Rheum* 2007;56:3928–39.
- 24 van der Heijde D, Klareskog L, Rodriguez-Valverde V *et al*. Comparison of etanercept and methotrexate, alone and combined, in the treatment of rheumatoid arthritis: two-year clinical and radiographic results from the TEMPO study, a double-blind, randomized trial. *Arthritis Rheum* 2006;54:1063–74.
- 25 Silman A, Symmons D, Scott DG, Griffiths I. British Society for Rheumatology Biologics Register. *Ann Rheum Dis* 2003;62(Suppl 2):ii28–9.
- 26 Joensuu JT, Huoponen S, Aaltonen KJ *et al*. The cost-effectiveness of biologics for the treatment of rheumatoid arthritis: a systematic review. *PLoS One* 2015;10:e0119683.
- 27 Stevenson M, Archer R, Tosh J *et al*. Adalimumab, etanercept, infliximab, certolizumab pegol, golimumab, tocilizumab and abatacept for the treatment of rheumatoid arthritis not previously treated with disease-modifying antirheumatic drugs and after the failure of conventional disease-modifying antirheumatic drugs only: systematic review and economic evaluation. *Health Technol Assess* 2016;20:1–610.
- 28 NICE. Guide to the Processes of Technology Appraisal (PMG19). 2014. www.nice.org.uk/process/pmg19 (date last accessed, 19 June 2017).
- 29 Chingcuenco F, Segal JB, Kim SC, Alexander GC. Bioequivalence of biosimilar tumor necrosis factor- α inhibitors compared with their reference biologics: a systematic review. *Ann Intern Med* 2016;165:565.
- 30 Smolen JS, Emery P, Fleischmann R *et al*. Adjustment of therapy in rheumatoid arthritis on the basis of achievement of stable low disease activity with adalimumab plus methotrexate or methotrexate alone: the randomised controlled OPTIMA trial. *Lancet* 2014;383:321–32.
- 31 Bijlsma JWW, Welsing PMJ, Woodworth TG *et al*. Early rheumatoid arthritis treated with tocilizumab, methotrexate, or their combination (U-Act-Early): a multicentre,

- randomised, double-blind, double-dummy, strategy trial. *Lancet* 2016;388:343–55.
- 32 Myasoedova E, Chandran A, Ilhan B *et al.* The role of rheumatoid arthritis (RA) flare and cumulative burden of RA severity in the risk of cardiovascular disease. *Ann Rheum Dis* 2016;75:560–5.
- 33 Bartlett SJ, Bykerk VP, Cooksey R *et al.* Feasibility and domain validation of rheumatoid arthritis (RA) flare core domain set: report of the OMERACT 2014 RA Flare Group Plenary. *J Rheumatol* 2015;42:2185–9.
- 34 Bingham CO 3rd, Pohl C, Woodworth TG *et al.* Developing a standardized definition for disease “flare” in rheumatoid arthritis (OMERACT 9 Special Interest Group). *J Rheumatol* 2009;36:2335–41.
- 35 Fautrel B, Morel J, Berthelot JM *et al.* Validation of FLARE-RA, a self-administered tool to detect recent or current rheumatoid arthritis flare. *Arthritis Rheumatol* 2017;69:309–19.
- 36 Choy EH, Smith CM, Farewell V *et al.* Factorial randomised controlled trial of glucocorticoids and combination disease modifying drugs in early rheumatoid arthritis. *Ann Rheum Dis* 2008;67:656–63.
- 37 Scott DL, Ibrahim F, Farewell V *et al.* Tumour necrosis factor inhibitors versus combination intensive therapy with conventional disease modifying anti-rheumatic drugs in established rheumatoid arthritis: TACIT non-inferiority randomised controlled trial. *BMJ* 2015;350:h1046.
- 38 Klarenbeek NB, van der Kooij SM, Guler-Yuksel M *et al.* Discontinuing treatment in patients with rheumatoid arthritis in sustained clinical remission: exploratory analyses from the BeSt study. *Ann Rheum Dis* 2011;70:315–9.
- 39 Rech J, Hueber AJ, Finzel S *et al.* Prediction of disease relapses by multibiomarker disease activity and autoantibody status in patients with rheumatoid arthritis on tapering DMARD treatment. *Ann Rheum Dis* 2016;75:1637–44.
- 40 Alivernini S, Peluso G, Fedele AL *et al.* Tapering and discontinuation of TNF-alpha blockers without disease relapse using ultrasonography as a tool to identify patients with rheumatoid arthritis in clinical and histological remission. *Arthritis Res Therapy* 2016;18:39.
- 41 Naredo E, Valor L, De la Torre I *et al.* Predictive value of Doppler ultrasound-detected synovitis in relation to failed tapering of biologic therapy in patients with rheumatoid arthritis. *Rheumatology* 2015;54:1408–14.